

JUN - 7 2001

K011370

APPENDIX M

510(k) SUMMARY

**SUMMARY OF THE SAFETY AND EFFECTIVENESS FOR POWDER FREE
PURPLE LATEX EXAMINATION GLOVES WITH & WITHOUT GRAPE SCENT**

Contact person : Cheah Chor Hee

This summary of safety and effectiveness information is being submitted in accordance with the requirements of SMDA 1990.

Device Information:

**Trade Name -- NON-STERILE POWDER FREE PURPLE LATEX EXAM
GLOVES
NON-STERILE POWDER FREE PURPLE LATEX EXAM
GLOVES WITH GRAPE SCENT**

Common Name - Exam gloves

Classification Name - Patient examination glove (per 21 CFR 880.6250)

Classification Information - Class I latex patient examination glove 80LYY, powder free and meeting all the requirements of ASTM-D3578-99 Standard Specification for Latex Examination Gloves for Medical Application.

Device Description:

Class I latex patient examination gloves 80LYY, powder free and meeting all the requirements of ASTM-D3578-99 Standard Specification for Latex Examination Gloves for Medical use.

Intended Use of Device:

A medical glove to be worn on the hand of the health care and similar personnel to prevent contamination between health care personnel and patient.

Technological Characteristics of Device:

1. Dimension

DIMENSION	ASTM D3578-99	SGMP
X-Small	70 mm +/- 10 mm	70 - 75 mm
Small	80 mm +/- 10mm	80 - 85 mm
Medium	95 mm +/- 10mm	90 - 97 mm
Large	111mm +/- 10mm	105 - 111 mm
Length	230 mm minimum for all sizes	245mm
Thickness - Finger Palm	0.08mm min 0.08mm min	0.08 mm min 0.08 mm min

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**2. Physical Properties (ASTM-D3578-99 Standard Specification for Latex Exam Gloves)
on Lot# 0070**

	TENSILE STRENGTH		ULTIMATE ELONGATION	
	ASTM-D3578-99	SGMP	ASTM-D3578-99	SGMP
Before Aging	Mpa	Mpa	%	%
	14.0		700	
X-Small		26.9		900
Small		27.2		860
Medium		29.6		910
Large		25.9		820
After Aging	14.0		500	
X-Small		24.4		910
Small		26.9		940
Medium		24.3		950
Large		23.2		830

3. Water Tight Test

Using the FDA specified 1,000 ml water leak test, 125 pieces of each size of the gloves were tested and our results are as given below:

BATCH #	SIZE	SAMPLE SIZE	LEAK STATUS	NUMBER LEAKED
UN-AGED				
0070	X-Small	125	Yes	2
0070	Small	125	Yes	1
0070	Medium	125	Yes	2
0070	Large	125	Yes	1
AGED				
0070	X-Small	125	No	0
0070	Small	125	Yes	1
0070	Medium	125	Yes	1
0070	Large	125	No	0

The above figures are within the draft FDA/ASTM D3578-99 requirements for latex exam gloves of 2.5% AQL.

4. Biocompatibility

The bio-compatibility test results are as per attached in APPENDIX L and show that the gloves passed the tests for examination gloves.

5. Total Residual Powder Content & Presence of Cornstarch

TESTS	FDA INTERNAL REQUIREMENT	SGMP's
Residual Powder Content (ASTM D 6124-00)	2 mg/glove max	Range: 0.4-0.9g/glove Mean : 0.6mg/glove
Presence of Cornstarch	Negative	Negative

6. Residual Protein Level

TESTS	FDA ALLOWABLE LEVEL	CLAIMED LEVEL
ASTM D 5712-95	< 50 µg/g	< 50 µg/g

Conclusion:-

The data presented indicate that the Non-sterile Powder Free Blue latex examination glove with and without bubblegum scent

1. meets/exceeds ASTM- D3578-99 Standard Specifications For Latex Examination Glove,
2. meets FDA pinhole requirements,
3. meets FDA claim criterion of a powder free glove.
4. meets the protein labeling claim level at <50 µg/g.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JUN - 7 2001

SGMP Company Limited
C/O Ms. Janna P. Tucker
Official Correspondent
Tucker & Associates
198 Avenue De La D' Emerald
Sparks, Nevada 89434

Re: K011370
Trade/Device Name: Non-Sterile Powder Free Purple
Latex Examination Glove with and without Grape Scent
And with Protein Labeling (50 micrograms or less)
Regulation Number: 880.6250
Regulatory Class: I
Product Code: LYY
Dated: May 2, 2001
Received: May 4, 2001

Dear Ms. Tucker:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

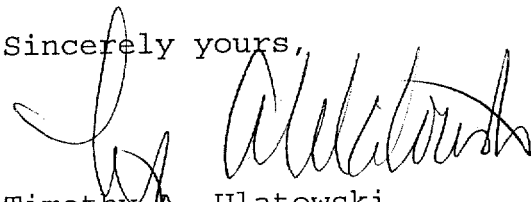
If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to

comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "<http://www.fda.gov/cdrh/dsma/dsmamain.html>".

Sincerely yours,



Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

K011370

INDICATION FOR USE STATEMENT

Applicant : SGMP Company Limited

510K NUMBER : K011370

Device Name : Non-sterile Powder Free Purple Latex Examination Gloves with & without Grape Scent + with Protein Labeling Claim (50mcgm or less) of total Extractable Protein per gram.

Indication For Use :

This is a medical glove to be worn on the hand of health care and similar personnel prevent contamination between health care personnel and the patient.

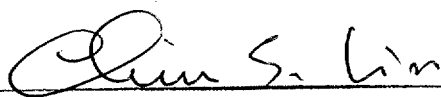
.....
Concurrence of CDRH Office of Device Evaluation (ODE)

**Prescription Use
Per 21 CFR 801.109**

OR

Over-The-Counter.....^X

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(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K011370